ORGANO PHOSPHATIC FERTILIZER

TECHNICAL FIELD

The present invention relates to a pelletized organic mineral fertilizer that comprises at least 40 percent (w/w) of dehydrated and biotreated pig manure and up to 60 percent (w/w) of a mineral fertilizer. The present invention also relates to a method for preparing a the pig manure-based organo phosphatic fertilizer of the present invention.

BACKGROUND OF THE INVENTION

The key macronutrients of a fertilizer are nitrogen (N), potassium (K) and phosporus (P). Although mineral fertilizers are good sources of compounds to provide a soil with those macronutrient, their efficacy is time limited. Indeed, in the immediate vicinity of a fertilizer pellet or granule of mineral fertilizer incorporated into the soil, the phosphorus fertilizer dissolved in a soil solution rapidly reacts with Fe, Al, or Ca compounds, hence decreasing fertilizer P availability to crops. In particular, phosphorus is strongly retained in podzolized soils used for potato production. Therefore, fertilizers able to slowly and gradually release P gained increasing interest over the last decades since they contribute to reduce the amount of fertilizer spread onto crops while maintaining a proper concentration of phosporus available to plants. The acidity or alcalinity of a soil also contributes to reduce the availability of P to plants and a pH ranging from 5.5 to 7.0 likely contributes to make P available.

The prior art reports that a combination of mineral and organic materials shows synergistic effects on the prevention of phosphorus binding by Fe, Al or Ca. P fixation is reduced since organic materials contain functional groups such as $-OH$, $-COOH$ and $-SO_3H$, that compete with orthophosphate ions for sorption sites in soils, thereby reducing phosphorus retention by Fe, Al or Ca. As manure and plant residues are good sources of organic materials, their combination with mineral phosphorus, or their application prior the application of a mineral fertilizer were shown to enhance P solubility. Among organic materials effective against soil retention of P, organic acids and humic substances are particularly effective in preventing P precipitation by Al compounds. The reactive organic ligands are bi- and tri-carboxylic acids as well as high molecular-weight humic and fulvic acids.

Although the application of organic matter may represent an interesting strategy to reduce P-fixation by Al, Fe or Ca ions, it does not alleviate reduced P availability attributed to soil pH conditions. For example, cow manure comprises a significant amount of alcaline cations that contribute to increase soil pH, which must be compensated by the application of lime. At the opposite, peat has a very low pH (around 4.3). The acidic condition of this organic matter may lead to a decrease of its efficiency as fertilizer since plant roots tend to avoid acidic conditions. Since it represents a very good organic source for its relatively high humic acid content, prior art reports the ammoniation of peat by associating it to a nitrogen source and equilibrating it into a potassium chloride solution, so as to increase the pH and the presence of ammonium humates, therefore providing a slow nitrogen-release fertilize. Considering the state of the prior art, it become highly desirable to be provided with an organic matter having excellent soil neutralizing and buffering properties without the need of additional supplementation or treatment.

Among the known organic matter sources usable for the fabrication of an organic mineral fertilizer, pig manure represents one of the most interesting alternative. Indeed, phosphorus in pig manure has been associated for many years with ground water and surface water contamination. The potential for water resource contamination by phosphorus therefore requires the implementation of regional management of animal manures and redistribution of excess nutrients. Pelletization of pig manure produces a dry and light-weight added-value commercial material that is easy to handle, transport, and apply and thus contributes to alleviate management problems. However, before drying and pelletizing, the manure must be liquid-solid separated to concentrate the solids, then bio-treated to eliminate odors. Since bio-treatment and drying of manure lead to additional expense, the latter must be compensated by sale of the pelletized manure. The nutrient composition of pelletized manure alone being relatively low compared to mineral fertilizers, its commercialization is modest since the nutrient composition is the main contributor of the market value of a fertilizer. In addition to its poor nutrient capacity, a pellet made from manure alone may also be too light-weight for bulk blending with mineral fertilizers.

International patent publication WO 02/0618 reports a method for producing an organic mineral fertilizer that comprises an organic material, such as biotreated pig manure, and urea are the nitrogen source. Although ammoniation of pig manure can occur

due to the presence of urea, the general purpose of this organic fertilizer is most likely to provide a soil with nitrogen since urea is the principal component of the fertilizer. Indeed, the organic material is restricted to a 30% w/w content. Therefore, such an organic mineral fertilizer is exposed to have a macronutrient content that is inadequate to respect the industry standard of a certified organo phosphatic fertilizer. Moreover, since the needs for pig manure are relatively low for the production of this fertilizer, it does not represent a particularly good way to alleviate the pig manure management problem that has been encountered for many years. Finally, the high nitrogen content of the organic fertilizer reported is not adequate for crops that have high phosphorus needs, such as corn and potatoes.

Considering the state of the prior art, there is a need for an organo-phosphatic fertilizer capable of solving pig manure management problems and having excellent soil neutralizing and buffering properties.

SUMMARY OF THE INVENTION

The present invention relates to an organo phosphatic fertilizer that comprises 40% to 90% of treated pig manure and 10 to 60% of a mineral fertilizer. The present invention also relates to a method for managing pig manure. The method of the present invention comprises sequentially or concomitantly treating and dehydrating the pig manure, mixing the treated and dehydrated pig manure with a mineral fertilizer in a proportion of 40% to 90% of pig manure for 10 to 60% of mineral fertilizer and pelletizing the mixture obtained therefrom.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1a to 1c are curves representing pH changes as a function of organic material concentrations in water.

Fig. 2 is a series of curves comparing the pH and buffering properties for different organic materials.

Fig.3a to 3c are curves showing variations in the concentration of soluble phosphorus as a function of organic material concentrations in water.

Fig. 4 represents a phosphorus partitioning flowchart and pool designation.

Fig.5 shows the increase in loosely bound P [$\Delta(LBP)$] due to added P as related to soil groups and addition of dry swine manure (LIOR), wherein r(LBP) is $\Delta Y/\Delta X$ from the origin point of the curve.

Fig.6 shows the increase in Al-sorbed P [$\Delta(SP_{Al})$] due to added P as related to soil groups and addition of dry swine manure (LIOR), wherein r(SP_{Al}) is $\Delta Y/\Delta X$ from the origin point of the curve.

Fig.7 shows the increase in Fe-sorbed P [$\Delta(SP_{Fe})$] due to added P as related to soil groups, wherein r(SP_{Fe}) is $\Delta Y/\Delta X$ from the origin point of the curve.

Fig. 8 shows the increase in organic P [$\Delta(P_{org})$] due to added P as related to soil groups and addition of lime or dry swine manure (LIOR).

Fig.9 shows the increase in desorbed P [$\Delta(DP)$] due to added P as related to soil groups and addition of dry swine manure (LIOR)n wherein r(DP) is $\Delta Y/\Delta X$ from the origin point of the curve.

Fig.10 is a flowchart illustrating the quantification of phosphorus partitioning for LSOM (minus LIOR and HSOM plus LIOR (highest P treatment) of treatments.

Fig. 11 is a curve showing corn yield as a function of LIOR concentration.

Figs. 12a and 12b are curves showing potato tuber yields as a function of P_2O_5/ha and LIOR concentration.

Fig. 13 is a curve showing the average of potato tubers with a diameter larger than 57 mm as a function of P_2O_5/ha.

Fig. 14 is a curve showing the soy grain yield as a function of LIOR concentration.

MODES OF CARRYING OUT THE INVENTION

The present invention now will be described more fully hereinafter with reference to the accompanying drawings, in which preferred embodiments of the invention are shown. This invention, may, however, be embodied in many different forms and

should not be construed as limited to the embodiments set forth herein; rather, these embodiments are provided so that this disclosure will be thorough and complete, and will fully convey the scope of the invention to those skilled in the art.

The present invention relates to a pig manure-based organo phosphatic fertilizer that comprises 40% to 90% (w/w) of treated pig manure and 10% to 60% (w/w) of a mineral fertilizer, and more preferably 50% to 80% (w/w) of said treated pig manure and 20% to 50% (w/w) of a mineral fertilizer. These proportions are preferred since they could contribute to significantly alleviate the problems related to management of pig manure while producing a fertilizer rich in phosphorus and capable of meeting the N-P-K requirements to be certified as an organo phosphatic fertilizer. Use of pig manure instead of another organic source alleviate the problem of pig manure management. By incorporating high concentrations of pig manure in the organic mineral fertilizer of the present invention, it facilitates the exportation of pig manure from region of high pig plant concentration to exterior zones. The method of the present invention therefore contribute to reduce the environmental drawbacks of phosphate over fertilization encountered for many years.

As used in the present description and claims, the term "treated pig manure" means that it has undergone a treatment known to those skilled in the art to stabilize it and make it odorless. The treated pig manure of the present invention may be obtained by aerobic treatment, anaerobic treatment, biofiltration, composting chemical treatment, thermal treatment or physico-chemical treatment. The purpose of treating the pig manure prior its use in the manufacture of a fertilizer is mainly to stabilize it and to makes it odorless. The treated pig manure may optionally be dehydrated prior to being mixed with the mineral fertilizer. However, treatment of the manure should not comprise the incorporation of structuring matter such as bark, for example, since it may prevent a proper pelletization of the organo phosphatic fertilizer.

The organo phosphatic fertilizer of the present invention is preferably a solid fertilizer and more preferably a pellet, a granule, a powder or a crumb. The pig manure is a high density organic matter and therefore, is easier to pelletize than low density matter, such as peat for example. In a preferred embodiment of the present invention, the organo phosphatic fertilizer may further comprise a binding agent that enhance pelletization. The

binding agent may comprise a zeolite, a silica, an attapulgite clay, a bentonite, or a polymer.

For example, the binding agent may be Cal-Ben™, Microsorb® LVM, Microsorb® RVM and Attagel®. Min-U-Gel® 200 is however preferred. Min-U-Gel® 200 is an attapulgite clay provided by ITC Minerals & Chemicals and available from Fluoridin, Inc. This product is currently used for the pelletization of a fertilizer including a binding agent for chicken and bovine manure. The binding agent is preferably added at a concentration ranging from 0% to 5% (w/w) and more preferably from 0.5% to 2% (w/w), so as to provide a proper binding of the organo phospatic fertilizer components while respecting the economical aspect of a fertilizer intended to be used on large surfaces. The physical and mechanical characteristics of the organo phosphatic fertilizer of the present invention are preferentially similar to those of known chemical fertilizers, making it easy to transport, store and spread.

The mineral fertilizer used for the purpose of the present invention may be any proper mineral fertilizer, but is preferably urea, monoammonium phosphate (MAP), diammonium phosphate (DAP), ammonia, magnesium sulfate, magnesium chloride, magnesium silicate, dolomite or chrysotyle. The presence of such fertilizers may contribute to the ammoniation of the organic matter and enhances the presence of soluble carbon and of slow-release P sources, such as struvite ($NH_4MgPO_4 \cdot 6H_2O$).

All patents, patent applications, articles and publications mentioned herein, both supra and infra, are hereby incorporated herein by reference.

The present invention will be more readily understood by referring to the following examples which are given to illustrate the invention rather than to limit its scope.

EXAMPLE I

Properties of different sources of organic material.

The prior art reports the combination of an organic material, such as peat, with a mineral fertilizer such as DAP and MAP, to create a synergy within a confined micro-environment rich in organic matter (OM) and favorable to the nutrition of plants. Mineral fertilizers have a wide range of pH and their combination with an organic material may significantly affect the pH of the resulting OMF. Indeed, some mineral fertilizer are acidic such as MCP (pH: 1.48), MAP (pH: 3.47) and MKP (pH: 3.99) while DAP (pH: 7.98) and DKP (pH: 10.1) are alkaline. The characteristics and fertilizing capabilities of the different organic sources remaining undetermined, we performed a systematic characterization of the mineral properties and binding, neutralizing and salinizing capabilities for dehydrated and biotreated pig manure (LIOR), composted bovine manure (CBV), composted chicken manure (CCM), composted sheep manure (CSM), composted leaves, bark and grass (CLBG), composted paper plant sludges (CPPS), peat and ammoniated peat (AP) Table 1).

Table 1

Origin of the tested organic matters.

Organic Matter	Source
Dehydrated and biotreated pig manure	DE.C Technologies, Inc. Québec, Canada
Composted bovine manure	Les composts Fafard, Inc. Québec, Canada
Composted chicken manure	Les composts Fafard, Inc. Québec, Canada
Composted sheep manure	Les composts du Québec, Inc. Québec, Canada
Composted leaves, bark and grass	Les composts du Québec, Inc. Québec, Canada
Composted paper plant sludges	Les composts du Québec, Inc. Québec, Canada
Peat	Premier Tech, Inc. St-Henri, Canada
Ammoniated peat[1]	Premier Tech, Inc. St-Henri, Canada

[1]Ammoniated peat was obtained by treating peat with NH_4OH 14.53N, in a proportion of 30 ml NH_4OH for 114 g of peat, according to Abbès *et al.*, 1994 and U.S. Patent Number 5,749,934.

pH Determination

Since any method designated for the purpose of determining the pH value of organic fertilizers were known, methods known in the art for pH determination of mineral fertilizer pH were adapted accordingly to the mineral saturation of each source of organic material. Indeed, mineral saturation of mineral fertilizer is easily achieved by observing the presence of unsolubilized material. Due to the particular nature of organic fertilizers, several of their components cannot be solubilized therefore avoiding the determination of saturation by the observance of precipitates.

To determine the organic matter/water ratio where the solutions pH was saturated, increasing amounts of every dried organic matter (see Table 2) were mixed with 70 mL of water in 250 mL polypropylene recipients, in duplicate, and agitated for forty-eight (48) hours at 300 rpm on a New-Brunswick type agitator. The amount of organic material mixed with water was determined according to the density of each organic material since it influences water retention and thus water available for further analyses. Samples were centrifuged at 10,409 x g for 10 minutes and filtered on a 2.5 μm number 42 Whatman paper (Fisher Scientific, Nepean, ON, Canada). The pH determination was done as reported in the prior art for each sample.

Table 2

Density of the tested organic matter.

	Organic Matter	Density (Mg/m^3)	Amount (g/ 70mL of water)
Low Density	Peat	0.20	0, 3, 6, 9, 12, 15, 18
	AP	0.25	0, 3, 6, 9, 12, 15
Medium Density	CSM	0.29	0, 3, 6, 9, 12, 15, 18, 21, 24
	CBM	0.30	0, 3, 6, 9, 12, 15, 18, 21, 24
	CCM	0.38	0, 3, 6, 9, 12, 15, 18, 21, 24, 27, 30
High Density	CLBG	0.48	0, 5, 10, 15, 20, 25, 30, 35, 40
	CPPS	0.52	0, 5, 10, 15, 20, 25, 30, 35, 40
	LIOR	0.67	0, 5, 10, 15, 20, 25, 30, 35, 40

Results

The pH decreases with increasing amount of organic matter until it reaches a critical ratio organic matter/water, which is specific for each substrate (Fig. 1 and Table3). The acidification of the solution is attributable to increasing amounts of organic acids comprised in the organic matter. Interestingly, the pH increased with increasing amounts of CSM and CBM (Fig 1). Having reached the critical organic matter/water ratio, the pH with LIOR, CSM, CBM and AP were 6.33, 6.91, 7.34 and 7.73, respectively. The pH of these organic materials is therefore approximately neutral, at the opposite of peat which have an acidic pH of 4.13. Therefore, LIOR, CSM, CBM and AP would be more appropriate for the production of an organic mineral fertilizer than untreated peat.

Table 3
pH at saturation.

Organic Matter	Density (Mg/m)	pH	Organic Matter / Water ratio at saturation level (g per 70 ml of water)
LIOR	0.67	6.33	25
CPPS	0.52	7.99	9
CLBG	0.48	7.74	20
CCM	0.38	7.84	18
CBM	0.30	7.34	21
CSM	0.29	6.91	15
Peat	0.2	4.13	9
AP	0.25	7.73	9

Determination of buffering properties

To determine the buffering capacities of the different organic matters, known methods previously reported were adapted. Briefly, we placed 1 g of the different dried organic matters into a fifty (50) mL polypropylene centrifuge tube. Twenty-five (25) mL of a solution comprising 0, 5, 10, 15, 20 or 25 mL 0.0025M H_2SO_4 or 0.025M NaOH were added to the organic matter and the solutions were agitated for 22 hours at 300 rpm on a New-Brunswick type shaker (Fisher Scientific, Nepean, ON, Canada). Samples were centrifuged for ten (10) minutes at 38,724 g. Supernatants were filtered on a <8μm

Whatman™ number 40 paper (Fisher Scientific, Nepean, ON, Canada) and the pH was monitored as reported in the art.

Results

Results show that LIOR has the most stable pH and therefore, has the best buffering capacities (Fig. 2). According to this method LIOR has a pH of 6.98, which is the closest to neutrality. LIOR could therefore be used with any acidic or alkaline mineral fertilizer. Peat has a lower pH and therefore is less adequate to be used as organic matter component in a organic mineral fertilizer, unless being treated to increase its alkalinity.

Physico-Chemical properties of organic matters

Each organic matter was treated with perchloric and nitric acids, according to Barhnisel and Bertsch (A.L. Page et al. eds., Methods in soil analysis. Part 2. Soil Sci. Soc. Am. Book Ser. 5, Madison, WI, 1982, pp 279-282). Phosporus was quantified according to the vanado-molybdate method. The different components of organic matter were quantified as known in the prior art.

Results

Table 4 shows the components of the different organic matters. Results of the physico-chemical analysis underline that LIOR has the highest content in macronutrients, namely nitrogen, phosphorus and potassium, with concentrations of 34.9, 41.5 and 72.8 g/kg of biomass, respectively. Particularly, the phosphorus concentration is nearly 2-folds higher in LIOR than in CCM, which is the second most concentrated organic matter in phosphorus.

Saturation in soluble organic carbon (SOC)

The method for determining the saturation in soluble organic carbon of the different organic matters was performed as described for the determination of pH saturation. The measurement of SOC is known in the art.

Table 4

Chemical properties of the tested biosolids.

	Peat	LIOR	CBM	CCM	CSM	CLBG	CPPS	AP
				g kg^{-1}				
M.O.	986	393	757	683	728	489	425	955
M.S.	918	923	818	783	934	940	953	878
C	572	228	439	396	422	284	247	554
Ntotal	15.8	34.9	19.5	32.2	18.4	14.2	6.5	30.9
NH$_4$-N	14.5	2.8	2.5	30.3	0.8	11.2	0.4	29.1
NO$_3$-N	1.3	32.1	17.0	1.9	17.6	3.0	6.1	1.8
P	0.6	41.5	2.4	20.7	7.4	2.4	1.4	0.04
K	0.1	72.8	7.2	12.8	11.0	9.2	2.0	0.1
Ca	0.0	17.9	22	10.1	13.9	13.4	17.9	2.6
Mg	0.2	20.1	2.6	5.0	5.1	3.7	2.6	0.5
S	0.7	0.8	3.6	1.8	2.9	0.6	0.4	2.1
Na	0.03	32.37	3.11	4.02	2.60	0.26	1.10	0.15
				mg kg^{-1}				
Al (ppm)	1396	973	2532	2999	4958	9689	23356	923
Fe (ppm)	567	6235	2089	3016	5669	8559	7613	684
Mn (ppm)	15	999	101	289	420	579	295	15
Cu (ppm)	10	912	19	99	143	31	130	4
Zn (ppm)	57	2040	69	270	376	144	275	37
Cr (ppm)	3.5	165.3	4.3	5.2	8.9	13.2	6.2	0.4
Pb (ppm)	1.1	0.3	1.4	1.9	3.4	20.7	31.7	0.3
Mo (ppm)	0.4	35.2	1.4	2.9	2.3	0.7	2.4	0.1
Co (ppm)	0.6	7.9	1.2	1.7	3.3	4.1	5.0	0.2
Ni (ppm)	4.0	143	6	10	12	10	14	1
As (ppm)	0.9	2.1	3.5	8.7	2.0	3.8	2.9	0.8
Cd (ppm)	0.2	0.7	0.1	0.3	0.3	0.5	0.8	0.1
				Without unit				
pH water	4.13	6.33	7.34	7.84	6.91	7.74	8.00	7.73
				dS m^{-1}				
CE	0.17	20.19	6.68	11.04	8.25	3.95	2.58	2.58
Category	C1*P1**	C2PI	C1P1	C1P1	C2P1	C1P1	C2P1	C1P1

C1 means any restriction on the use of the biosolid regarding to its content in inorganic contaminants.
C2 means some restriction on the use of the biosolid since some inorganic contaminants are over environmental standards.
P1 means any restriction on the use of the biosolid regarding to pathogen contaminants.

Results

Table 5 shows the organic matter/water ratio corresponding to saturation in SOC and the corresponding SOC concentration. The highest ratio was obtained with LIOR, with 9,900 mg/L, followed by chicken manure (6,800 mg/L) and ammoniated peat (approximately 6,300 mg/L). LIOR therefore represents an excellent source of binding agent that contributes to increase the availability of phosphorus within the micro-environment of a pellet, by competing with Al, Fe and Ca. Since LIOR has the highest

SOC ratio, it is likely the best competitor of Fe, AL and Ca for phosphorus binding in a soil.

Table 5

Soluble Organic Carbon (SOC) concentration at saturation

Organic Matter	Density (Mg/m^3)	Concentration of SOC at the plateau (mg/L)	Biosolid / water ratio at saturation (g per 70 ml of water)
LIOR	0.67	9,900	25
CBP	0.52	165	23
CFEG	0.48	780	25
Cpoule	0.38	6,800	18.5
CB	0.30	660	11
CM	0.29	180	10
Tourbe	0.2	2,110	12
TA	0.25	> 6,300*	--

* The SOC plateau has not been reached with TA. The indicated SOC concentration corresponds to a Biosolid/Water ratio of 15 g per 70 ml of water.

EXAMPLE II

Quantification of soluble phosphorus.

Soluble phosphorus quantification was determined as known in the art. Results illustrated on Figs. 3a to 3c are expressed as concentration of soluble phosphrus (mg/L) as a function of the amount of organic matter (g per 70 mL of water). Table 6 shows that LIOR has the highest concentration in soluble P (590 mg/L), nearly 300-time more elevated than peat (2 mg/L). Therefore, LIOR is the most appropriate choice for the manufacture of an organo phosphatic mineral since it contributes to reduce the cost attributed to the phosphatic fertilizer portion of the pellet while maintaining a proper amount of phosphorus within the organo phosphatic fertilizer.

Table 6

Soluble phosphorus concentration at saturation.

Biosolid	Density (Mg m^{-3})	Soluble P concentration mg L^{-1}	Biosolid / Water ratio (g par 70 ml)
LIOR	0.67	590	13.2
CPPS	0.52	1.2	15
CLBG	0.48	20.3	25
CCM	0.38	342	15.9
CBM	0.30	38	10.6
CSM	0.29	52	15
Peat	0.2	2.0	9.9
AP	0.25	18	15

EXAMPLE III

Determination of pig manure enrichment on phosphorus transformation is acid-light-textured soils.

Materials and Methods

Soil and Manure Analyses

Four surface soil samples (0-20 cm) were collected from fluvio-glacial or deltaic deposits in St-Ubalde, Quebec, Canada, where potato (*Solanum tuberosum* L.) and small grains are grown in rotation. Three humo-ferric podzols (Morin and Bevin sandy loams, Ivry loamy sand) were low in SOM (LSOM), and an Ivry peaty phase was high in organic matter content (HSOM). Soil samples were collected in the potato phase.

Soil samples were dried at 105°C and passed through a 2-mm sieve. Soil pH was measured in 0.01 M CaCl$_2$ using a 1:1 soil to solution ratio. Organic C was determined by the Walkley-Black procedure (Nelson and Sommers, A.L. Pages et al. eds., Methods of soil analysis. Par 2. Agronomy Monogr. 9, Am. Soc. Agronom. , Madison, WI, 1982, pp 539-579). Soil texture was analyzed by the hydrometer method (Day, P.R., C.A. Black eds., Methods of soil analysis. Part 1. Physical and mineralogical properties. Am. Soc. Agron., Madison, WI. 1965, pp545-567). The P and Al were extracted using the Mehlich-III procedure (Mehlich, A., 1984, Commun. Soil Sci. Plant Anal. 15:1409-1416). Phosphorus was determined colorimetrically (Laverty, J.C., 1963, Soil Sci. Soc. Am. Proc.

27:360-361), and Al by atomic absorption spectrophotometry (AAS) (Perkin Elmer 603 spectrophotometer, Perkin Elmer, Wellesley, MA). The $100(P/Al)_{M-III}$ weight ratio is a measure of soil P saturation for soil fertility classification and environmental risk assessment (Khiari et al., 2000, Environ. Qual., 29:1561-1567). Soils with a $100(P/Al)_{M-III}$ weight ratio between 2 and 4% are considered to be of very low P fertility level (low P availability) and at very low P environmental risk (high P fixation) (Khiari et al., 2000, Environ. Qual., 29:1561-1567).

The oxalate and pyrophosphate extractions were performed according to McKeague (1978). Soil samples were ground to <150 μm-sieved. The mixtures were 2.5-□m gravity-filtered (Whatman no. 42 paper), and the filtrate analyzed by plasma emission spectroscopy. The acid ammonium oxalate extracts organically bound, amorphous, and some crystalline forms of Al and Fe. The pyrophosphate extracts mainly organically bound, and only very small amounts of other forms of Al and Fe. The degree of phosphorus saturation (DPS) was computed as follows (Breeuwsma and Silva , 1992, Agric. Res. Dep. Rep. 57, Winand Staring Centre for Integrated Land, Soil, Water Res., Wageningen, The Netherlands):

$$DPS(\%) = \frac{100 P_{ox}}{\alpha_m (Al_{ox} + Fe_{ox})} \qquad (1)$$

where P_{ox}, Al_{ox} and Fe_{ox} are oxalate-extracted P, Al and Fe; DPS is the degree of phosphorus saturation expressed on a molar basis; α_m, the maximum saturation factor for total sorption, is equal to approximately 0.66 across a wide variety of soils (Khiari et al., 2000, Environ. Qual., 29:1561-1567). Lime addition as reagent-grade $CaCO_3$ was based on buffer pH (Shoemaker et al. 1961, Soil. Sci. Soc. Am. Proc. 25:274-277) to achieve a pH of 6.5 in the soil volume. The LIOR contained 368 g total C kg^{-1}, 26 g soluble C kg^{-1}, and 23.3 g total P kg^{-1}. Total C was determined by combustion. Soluble C was extracted in a saturated solution of 30 g LIOR in 70 ml of distilled water. The mixture was shaken for 24 h on an end-over-end shaker at 300 rpm, centrifuged at 12 000 rpm, then gravity-filtered through a Whatman no. 42 paper. Soluble C was digested according to Nelson and

Sommers (1982). Total P was obtained after digesting LIOR in a HNO_3-$HClO_4$ mixture (Barnhisel and Bertsch, A.L. Pages, eds., Methods of soil analysis. Part 2 Agronomu Monogr. 9, Am. Soc. Agron., Madison, WI., 1982, pp539-579). Total phosphorus was quantified using the yellow method (Kuo, S., 1996 Phosphorus, Pages 869-919, in Soil Sci. Soc. Am. Book, Ser. 5. Methods of soil analysis, Madison, WI).

Sequential P Fractionation

The procedure for determining P pools and their designation are given in Table 7. The sequential fractionation of designated inorganic P (P_i) pools was conducted using a modified Chang-Jackson procedure as described and designated by Kuo, S., (1996, Phosphorus, Pages 869-919, in Soil Sci. Soc. Am. Book, Ser.5 . Methods of soil analysis, Madison, WI). All extracts were 2.5-μm gravity-filtered (Whatman no. 42 paper). The loosely bound P_i (LBP) was extracted using 1.0 M NH_4Cl, the P sorbed by Al (SP_{Al}) using 0.5 M NH_4F, the P sorbed by Fe (SP_{Fe}) using 0.1 M NaOH, the P sorbed by Ca (SP_{Ca}) using 0.25 M H_2SO_4, and the P sorbed as occluded or reductant P_i (SP_{red}) using a citrate-dithionite-bicarbonate solution. All fractions except SP_{red} were determined according to (Kuo, S., 1996, Phosphorus, Pages 869-919, in Soil Sci. Soc. Am. Book, Ser. 5. Methods of soil analysis, Madison, WI), SP_{red} according to Peterson and Corey (1966).

Separate P Analyses

Total P was determined after digesting the soil (< 2 mm) in a HNO_3-$HClO_4$ mixture. Soil organic P was quantified separately using a basic EDTA procedure (Bowman and Moir, 1993, Soil Sci. Soc. Am. J. 57:1516-1518) and 0.5 g soil samples. Samples were extracted for 2 h at 85°C with 25 ml of 0.25 M NaOH + 0.05 M Na_2EDTA. Organic P (P_{org}) in extracts was determined by persulfate oxidation. The P recovery was computed as the sum of inorganic and organic P fractions divided by total P. The P recovery (mean ± standard deviation) was 97.1 ± 3.7% across soils and treatments. The phosphorus was quantified using the yellow method (Kuo, S., 1996, Phosphorus, Pages 869-919, in Soil Sci. Soc. Am. Book, Ser. 5. Methods of soil analysis, Madison, WI). The desorbed P pool (DP) was determined in separate subsamples using 1:60 water to soil volume ratio and filtered (< 2.5 μm, Whatman™ no. 42 paper). The P was quantified by the ascorbic acid blue method (Kuo, S. 1996, Phosphorus, Pages 869-919, in Soil Sci. Soc. Am. Book, Ser. 5. Methods of soil analysis, Madison, WI).

Table 7

Methodology for designated P pools.

P pool	Symbol	Extraction procedure or computation method	
		Determined by extraction on the same sample	
Loosely bound P pool	LBP	$1\,M\,NH_4Cl$ (Kuo, 1996)	
Al-sorbed P pool	SP_{Al}	$0.5\,M\,NH_4F$ (Kuo, 1996)	
Fe-sorbed P pool	SP_{Fe}	$0.1\,M\,NaOH$ (Kuo, 1996)	
Ca-sorbed P pool	SP_{Ca}	$0.25\,M\,H_2SO_4$ (Kuo, 1996)	
Reductant P pool	SP_{red}	Citrate-dithionite-bicarbonate (Kuo, 1996)	
		Determined by extraction on separate samples	
Desorbed P Pool	DP	Sissingh (1971) and Van der Zee et al. (1987)	
Organically converted P	P_{org}	$0.05\,Na_2EDTA + 0.25\,NaOH$ (Bowman and Moir, 1993)	
		Deduced by computation	
Sorbed inorganic P pool	SP_{inorg}	$SP_{Al} + SP_{Fe} + SP_{Ca} + SP_{red}$	(Eq. 6)
Reversible adsorbed P pool	AP_{rev}	$DP - LBP$	(Eq. 5)
Sorbed P pool due to slow reaction	SP_{sr}	$SP_{inorg} - AP_{rev}$	(Eq. 7)

Simulated P Diffusion Volume

The P diffusion coefficient from fertilizer is small in the range of 5 to $23.710^7\ cm^2\ s^{-1}$, depending on P rate, soil water content, and bulk density (BD) (Hira and Singh, 1978, Soil Sci. Soc. Am. J. 42:561-565). Hira and Singh (1977, Soil Sci. Soc. Am. J. 41:537-540) found that the P diffusion volume in soils increased with moisture content and BD, with maximum at $1.60\ Mg\ m^{-3}$ followed by a drop toward $1.75\ Mg\ m^{-3}$. Moisture content was found to be $0.15\text{-}0.18\ m^3\ m^{-3}$ in a Haibowal silty clay loam and $0.18\text{-}0.25\ m^3$ m^{-3} in a Choa sandy loam for maximum ^{36}Cl diffusion with BD of $1.25\ Mg\ m^{-3}$. It is knpwn that molecular diffusion coefficient depended on soil moisture content and BD. Similar coefficients were obtained for a fine sand (BD = $1.68\ Mg\ m^{-3}$) and a loam (BD = $1.52\ Mg\ m^{-3}$) with a volumetric water content of $0.25\ m^3\ m^{-3}$, and for a peat-perlite mixture (BD = $0.15\ Mg\ m^{-3}$) with a volumetric water content close to $0.37\ m^3\ m^{-3}$ (Riga and Charpentier, 1998). Our LSOM soils had BD values (scooped soil sample) between 1.17 and $1.36\ Mg\ m^{-3}$, compared to $0.78\ Mg\ m^{-3}$ for the HSOM soil. Moisture contents were adjusted to $0.20\text{-}0.25\ m^3\ m^{-3}$ in the LSOM soils, and to $0.37\ m^3\ m^{-3}$ in the HSOM soil for facilitating molecular diffusion into the prescribed diffusion volume.

Dissolved P diffuses away from the granule across a soil volume of two to three times the diameter of the granule . The P distribution pattern from mono-ammonium phosphate (MAP) in an acid Hartsells fine sandy loam showed a P diffusion diameter of 38, 40 and 41 mm, respectively, around the application point after 4, 14 and 49 days. Should the diffusion volume be described by spheres of 20.5 mm of radius contacting each other along a continuous line and should the potato row spacing be 0.915 m, the total diffusion volume of two fertilizer bands about the potato seed would be about 19 m^3 ha^{-1}. The simulated P diffusion volume was thus computed as follows:

$$V = (R^3 - r^3)/r^3 \qquad (2)$$

where V is the simulated diffusion volume (35 ml); R, the radius of the external limit of the P diffusion sphere after 6 wk of incubation, was set equal to 20.5 mm; r was set equal to R/3, giving a spherical volume of 1.35 ml. Thus, the selected granule to soil volume ratio was 1 to 26 (35/1.35).

Soil Treatments

Lime, manure, and fertilizers were applied into a constant soil volume. Treatments are described in Table 8. A volume of 1.35-ml (800 mg) of LIOR was added to a 35-ml soil sample, i.e. 26 times LIOR volume, to provide 23 g LIOR L^{-1} of soil (= 800 mg of LIOR per 35 ml of soil) or 14.6 g L^{-1} of soil as organic matter (OM). Added OM was 21 g kg^{-1} for the Morin SL, 22 g kg^{-1} for the Bevin SL, 19 g kg^{-1} for the Ivry LS, and 32 g kg^{-1} for the Ivry LS, peaty phase, due to differences in BD. Adding up exogenous and indigenous OM percentages, OM contents were 61 g kg^{-1} in the Morin SL, 71 g kg^{-1} in the Bevin SL, 59 g kg^{-1} in the Ivry LS, and 232 g kg^{-1} in the Ivry LS peaty phase, respectively. The Morin and Bevin soils received 276 mg of $CaCO_3$ per 35 ml of soil, the Ivry soil 185 mg, and the Ivry peaty phase, 369 mg. Added P as MAP- [reagent-grade mono-ammonium phosphate: $NH_4H_2PO_4$] and LIOR-P was 0, 49, 127, or 265 mg P per 35 ml of soil. Added P divided by the weight of the MAP-LIOR mixture were 0 (zero-P control), 5, 10, and 15%. Due to variations in BD, the P added into a 20.5-mm radius fertilizer band was, on a weight basis for 265 mg P per 35 ml of soil, as follows: 6360 mg P kg^{-1} for the Morin SL,

6470 mg P kg^{-1} for the Bevin SL, 5570 mg P kg^{-1} for the Ivry LS, and 9706 mg P kg^{-1} for the Ivry LS peaty phase. Application rates would be respectively 0, 27, 69 and 144 kg P ha^{-1}, the lower rates being commonly applied to potato (*Solanum tuberosum* L.) grown in high-fertility soils, and the intermediate rate being applied in medium-fertility soils (Khiari et al. 2000). The potato could respond to the highest P rate in high-P fixing soils. There was no pre-incubation of soils and treatments, which were intended to modify soil properties in the fertilizer band. Prior to the experiment, dry samples of 35 ml of soil were mixed with MAP, lime or LIOR in 250 ml polypropylene recipients. The four soils, maintained at field capacity with distilled water, were incubated in duplicates at 23 ± 2°C for six wk to allow slow reactions to occur.

Table 8

Treatments applied to the incubated soils (MAP = mono-ammonium phosphate, and LIOR = dry swine manure)

Ligand	Treatment identification	MAP weight	MAP+LIOR weight [z]	Total P as MAP
		mg of product in 35 ml of soil		
	0 P	0	800	0
LIOR (800 mg per 35 ml of soils)	5 P	184	984	49
	10 P	470	1270	127
	15 P	980	1780	265
	0 P	0	0	0
CaCO$_3$ (185-369 mg per 35 ml of soils)	5 P	184	184	49
	10 P	470	470	127
	15 P	980	980	265
	0 P	0	0	0
Non-amended control	5 P	184	184	49
	10 P	470	470	127
	15 P	980	980	265

[z] Total weight of fertilizer granule = weight of MAP + LIOR, LIOR = 800 mg for the LIOR treatment and LIOR = 0 for others

Fertilizer Phosphorus Accumulation in Soil P Pools

Net P acquisition in a given P pool was made on a volume basis (mg L^{-1}). A unitless rate of P acquisition per unit of added P was computed for the prescribed diffusion

volume. Differential increase in P pool $\Delta(P)$ as net P acquisition in mg P L^{-1} was computed by difference $\Delta(P)_F$ between P pools in MAP-fertilized (P_F) and zero-P control (P_C) treatments, as follows:

$$\Delta(P) = P_F - P_C \tag{3}$$

Values $\Delta(LBP)$, $\Delta(SP_{Al})$, $\Delta(SP_{Fe})$, $\Delta(SP_{red})$, $\Delta(SP_{Ca})$., $\Delta(DP)$, and $\Delta(P_{org})$ were computed. Proportions $r(P)_F$ in a given pool relative to added P (mg P L^{-1}) was computed as follows:

$$r(P)_F(\%) = \frac{\Delta(P)_F}{Added\,P} \times 100 \tag{4}$$

The $r(P)_F$ was the slope of the relationship between $\Delta(P)$ and added P, computed similarly to the increase in anion exchange P pool in response to added P (Jones et al., 1984). Loosely bound P, designated as LBP (NH$_4$Cl-extracted), is a fraction of desorbed P. Difference between desorbed P and LBP was designated as reversibly adsorbed P ($r(AP_{rev})_F$) computed as follows:

$$r(AP_{rev})_F = r(DP)_F - r(LBP)_F \tag{5}$$

where $r(AP_{rev})_F$, $r(DP)_F$, $r(LBP)_F$ are proportions of reversibly adsorbed RP, desorbed RP (Sissingh 1971), and loosely bound RP pools, respectively. Proportions of sorbed inorganic P pools were summed up as $r(SP_{inorg})_F$ as follows:

$$r(SP_{inorg})_F = r(SP_{Al})_F + r(SP_{Fe})_F + r(SP_{red})_F + r(SP_{Ca})_F \quad (6)$$

Sorbed P due to slow reactions (Van der Zee et al., 1987), i.e. $r(SP_{sr})_F$, was computed by difference between sorbed inorganic P (Eq. 6), and reversibly adsorbed P (Eq. 5) as follows:

$$r(SP_{sr})_F = r(SP_{inorg})_F - r(AP_{rev})_F \quad (7)$$

The sum of all P pools from added P was computed as follows:

$$r(P_{total})_F = r(SP_{inorg})_F + r(P_{org})_F + r(LBP)_F \quad (8)$$

See the flowchart of P pools presented in Fig. 4.

Statistical Analyses

The experimental setup was a factorially arranged randomized complete block design with two replications. We used the GLM procedure for data analysis (SAS, 1990, SAS/STAT User's Guide. Version 6. 4th ed. Statistical Analysis System Institute, Cary, NC). Soil type and ligand sources (LIOR or lime) were considered as categorical variables, while the P application rates were analysed as continuous variables. Significance of differences between means was assessed using orthogonal contrasts. Regression analyses on the effects of P application rates were conducted using the Excel package (Microsoft™, 1997) and the REG procedure (SAS, 1990, SAS/STAT User's Guide. Version 6. 4th ed. Statistical Analysis System Institute, Cary, NC).

Results

Partitioning of Added P among Phosphorus Fractions in Fertilized Soils

Soil characteristics and P fractions are presented in Table 9. The SP_{Al} accounted for 37 to 43% of total P in LSOM soils, compared to 9% in the HSOM soil. The SP_{Fe} fraction ranked second. Soil types, P rates, and amendments (LIOR, lime) significantly influenced the partition of added P (Table 10). The P rate and contrast between LSOM and HSOM soils gave highest F values. There were significant treatment interactions for $\Delta(LBP)$, $\Delta(SP_{Al})$, $\Delta(SP_{Fe})$, $\Delta(DP)$, and $\Delta(P_{org})$. Linear and quadratic effects of P doses depended on soils and amendments. The contrasts were significant between sandy loams (Morin, Bevin) on the one hand, and between sandy loams and the loamy sand (Ivry), on the other. There was no significant effect of lime compared to control across P fractions, except for $\Delta(DP)$ and $\Delta(P_{org})$. We thus contrasted LIOR with (control + lime). The $\Delta(DP)$ pool was much larger than the $\Delta(LBP)$ (Table 11). Similarly to 1 M KCl, the 1 M NH$_4$Cl solution extracts exchangeable and some non-isotopically exchangeable Al^{3+}. Presumably, some NH$_4$Cl-exchangeable Al^{3+} reacted with ortho-phosphate to form aluminium phosphates, that lowered the $\Delta(LBP)$ pool and was recoverable as $\Delta(DP)$. The $r(P)_F$ for the highest P treatment are presented in Table 11 for the soil times amendment interaction. 79 to 92% of added P was sorbed as SP_{inorg} across treatments in LSOM soils. Comparatively, 51 to 61% of added P was sorbed as SP_{inorg} in the HSOM soil. The $r(LBP)_F$ pool was abundant in the HSOM soil only.

Table 9

Properties of the acid soils under study.

Property		Soil Series			
		Morin	Bevin	Ivry	Ivry peaty phase
Texture		Sandy loam	Sandy loam	Loamy sand	Loamy sand
			g kg^{-1}		
Organic matter content		40	49	40	200
Clay		65	65	56	48
Sand		723	524	876	734
			Unitless		
pH (CaCl$_2$)		4.8	4.3	5.2	3.7
SMP buffer pH		5.6	5.6	6.0	5.2
			Mg m^{-3}		
Bulk density		1.19	1.17	1.36	0.78
			mg kg^{-1}		
P total		2440	2645	1170	407
P Fractionation[z]	LBP	0	0	0	5
	SP$_{Al}$	1048	1126	431	37
	SP$_{Fe}$	545	655	159	18
	SP$_{red}$[y]	80	107	21	18
	SP$_{Ca}$	248	215	275	4
Al extracted by NH$_4$F		3591	5781	1056	397
Fe extracted by NaOH		244	162	153	700
P$_{M-III}$[x]		75.5	81.7	49.1	45.3
Al$_{M-III}$[x]		1888	1945	2046	1510
Fe$_{M-III}$[x]		206	191	144	658
			mmol kg^{-1}		
P$_{ox}$[w]		62.2	71.7	35.9	14.1
Al$_{ox}$[w]		404.4	474.1	351.1	129.5
Fe$_{ox}$[w]		167.1	182.9	107.9	109.3
Al$_{ox}$ + Fe$_{ox}$		571.5	657.0	459.0	238.8
			%		
DPS (%)[v]		16.5	16.5	11.8	8.9
100(P/Al)$_{M-III}$[u] (%)		4.0	4.2	2.4	3.0

[z] Differential P dissolution technique of Kuo (1996)
[y] Extracted by citrate-dithionite-bicarbonate
[x] Extracted using the Mehlich-III procedure (Mehlich, 1984)
[w] Oxalate extraction according to McKeague (1978)
[v] Degree of Phosphorus Saturation as defined by Breeuwsma and Silva (1992)
[u] P saturation as defined by Khiari et al. (2000)

Table 10

Effects of soil (LSOM = low soil organic matter; HSOM = high soil organic matter; SL = sandy loam; LS = loamy sand)), amendment (LIOR = dry swine manure), and P doses on P pools.

Source	Df[z]	P pool				
		Loosely bound P	*Sorbed P*			*Desorbed P*
		$\Delta(LBP)^y$	$\Delta(SP_{Al})^y$	$\Delta(SP_{Fe})^y$	$\Delta(P_{org})^y$	$\Delta(DP)^y$
		F value				
Soil	3	1156.56**	971.01**	300.20**	440.32**	450.41**
Amendment	2	27.76**	22.32**	0.36 ns	121.99**	16.58**
Dose P	2	2091.97**	10066.54**	721.77**	791.51**	2140.08**
Soil * Amendment	6	10.30**	1.05 ns	8.73**	12.11**	7.05**
Soil * dose P	6	580.70**	261.30**	29.41**	81.30**	248.05**
Amendment * dose P	4	17.18**	9.76**	0.30 ns	38.13**	6.89**
Soil * Amendment * dose P	12	8.67**	1.81 ns	1.30 ns	6.59**	4.75**
		Contrast				
LSOM vs HSOM	1	3456.56**	2911.05**	782.38**	1269.43**	1332.05**
LSOM (SL vs LS)	1	7.81**	1.48 ns	104.24**	48.32**	14.97**
Morin vs Bevin	1	5.31*	0.52 ns	13.97**	3.32 ns	4.20*
Control vs Lime	1	1.98 ns	1.68 ns	0.59 ns	36.59**	20.80**
(Control+Lime) vs LIOR	1	53.54**	42.97**	0.13 ns	207.40**	12.35**
		Polynomial contrast				
Linear (P rates)	1	4073.03**	20113.76**	1432.97**	1570.02**	4174.53**
Quadratic (P rates)	1	112.12**	18.21**	10.35**	12.75**	106.83**
Root of error mean of squares		45.32	98.20	53.90	25.00	131.32
Coefficient of variation		12.7%	3.3%	10.7%	13.5%	12.4%
R-Square		0.99	0.99	0.99	0.99	0.99

[z]df: degree of freedom
ns, *, **: non significant and significant at the 0.05 and the 0.01 levels, respectively
[y] Fertilizer phosphorus accumulation in soil P pool (Eq.[3])

Table 11
Effects of soil (LSOM = low soil organic matter; HSOM = high soil organic matter; SL = sandy loam; LS = loamy sand)), amendment (LIOR = dry swine manure), and P doses on proportions of added P in soil P pools for the highest P treatment

Source	Df[z]	P pool Loosely bound P $r(LBP)_F^y$	Sorbed P $r(SP_{Al})_F^y$	$r(SP_{Fe})_F^y$	$r(P_{org})_F^y$	Desorbed P $r(DP)_F^x$
		F value				
Soil	3	758.87**	669.48**	90.48**	176.88**	330.95**
Amendment	2	20.98**	19.41**	0.14 ns	66.65**	10.36**
Soil * Amendment	6	9.43**	157 ns	2.23ns	6.95**	5.82**
		Contrast				
LSOM vs HSOM	1	2270.04**	2004.64**	240.05**	501.44**	978.64**
LSOM (SL vs LS)	1	3.02ns	1.11 ns	31.30**	29.11**	11.26**
Morin vs Bevin	1	3.56ns	2.68 ns	0.09 ns	0.08 ns	2.95**
No ligand vs Lime	1	1.23 ns	2.68 ns	0.02 ns	20.09**	13.03**
(No ligand+Lime) vs LIOR	1	40.73**	36.14**	0.26 ns	113.20**	7.68**
Root of error mean of squares		1.03	1.81	1.16	0.53	2.89
Coefficient of variation		9.26%	2.67%	10.8%	12.0%	8.88%
R-Square		0.99	0.99	0.96	0.98	0.99

Average $r(P)_F$ across treatments

Soil	Amendment	$r(LBP)_F$	$r(SP_{Al})_F$	$r(SP_{Fe})_F$	$r(P_{org})_F$	(Total)[w]	$r(DP)_F$	$r(SP_{rev})_F^v$	$r(SP_{inorg})_F^u$	$r(SP_{irr})_F^t$
						%				
Morin	Control	4.9	78.8	9.6	1.8	**95.1**	22.0	17.1	88.4	71.3
	LIOR	6.1	74.3	10.0	4.1	**94.5**	22.6	16.5	84.3	67.8
	Lime	5.8	76.7	9.3	1.9	**93.7**	20.5	14.7	86.1	71.3
Bevin	Control	4.3	78.3	8.4	2.0	**93.0**	18.9	14.6	86.7	72.1
	LIOR	4.5	76.0	10.0	3.8	**94.4**	20.0	15.4	86.0	70.6
	Lime	4.5	80.7	11.1	1.7	**98.0**	17.6	13.1	91.8	78.7
Ivry	Control	5.3	80.5	6.5	2.4	**94.7**	25.4	20.2	87.0	66.8
	LIOR	7.2	72.7	6.2	5.0	**91.1**	25.2	17.9	78.9	60.9
	Lime	5.2	76.4	6.7	4.6	**93.0**	24.7	19.5	83.2	63.7
Ivry	Control	24.9	42.2	19.0	6.0	**92.0**	69.6	44.6	61.1	16.5
	LIOR	34.0	35.0	16.2	11.4	**96.6**	71.7	37.7	51.2	13.5
	Lime	26.1	40.0	15.4	8.6	**90.1**	52.3	26.1	55.3	29.2

[z] df: degree of freedom; ns, *, **: non significant and significant at the 0.05 and the 0.01 levels, respectively; [y] P proportion of added P in soil P pools (Eq.[4]); [x] P proportion of added P as desorbed P; [w] Sum of $r(LBP)_F$; $r(SP_{Al})_F$, $r(SP_{Fe})_F$, and $r(P_{org})_F$; [v] P proportion of added P as reversibly bound P (Eq.[5]); [u] P proportion of added P as sorbed inorganic P (Eq.[6]); [t] P proportion of added P as P sorbed by slow reactions (Eq.[7]).

Loosely Bound Phosphorus Pool (LBP)

The *Δ(LBP)* depended on soil type and amendment (Table 10). Only 0.7 to 6.1% of added P was converted to LBP in LSOM soils, compared to 34% in the LIOR-

treated HSOM soil (Fig. 5). The $\Delta(LBP)$ increased nearly 5 times as SOM increased from 4 to 20% (Table 9). The P fixation capacity was 2.4 times larger in LSOM soils [i.e. 0.66 times 562.5 mmol $(Fe_{ox}+Al_{ox})$ kg^{-1})] than in the HSOM soil [i.e. 0.66 times 238.8 mmol $(Fe_{ox}+Al_{ox})$ kg^{-1})], although DPS was low across soils (Table 9). Lower pH in the HSOM than in LSOM soils was expected to promote P fixation and thus decrease $\Delta(LBP)$ for comparable DPS values. However, ligand exchange reactions in the HSOM soil presumably protected added P by forming stable OM-metal complexes (Fox et al., 1990). Indeed, the pyrophosphate extraction accounted for 24.5% of the oxalate-extracted Al and Fe in the HSOM, compared to 16.5-21.3% in LSOM soils (Table 12). The LIOR increased pyrophasphate-extracted Al and Fe by 2.7% in the HSOM soil, and produced a significant effect only in the Ivry soil among LSOM soils (Table 12). The HSOM was by far the most responsive to LIOR in producing organically-bound Al and Fe (Table 12) and converting added P into LBP (Fig. 5). The DPS based on oxalate extraction alone (thus including the pyrophosphate extraction) should be interpreted with caution in connection with environmental protection and as a soil P availability index for crops, considering the apparent differential reactivity of amorphous and organically bound Al and Fe toward P.

Highest-level interaction and the (control + lime) vs LIOR contrast were significant for $\Delta(LBP)$ (Table 10), due to a significant LIOR effect in the HSOM soil at the highest P rate. The $r(LBP)_F$ (Eq.4) in the HSOM soil averaged nearly 25% across control and lime treatments, and 34% with LIOR (Table 11). In LSOM soils, LIOR showed no significant effect compared to (control + lime) for the Bevin soil, but a slight contribution to LBP of less than 2% in the Morin and the Ivry soils (Fig. 6). Higher amounts of LIOR than used in this study should likely be added to increase the organic amendment effect in those LSOM high P-fixing soils.

Phosphorus Sorbed by Aluminium (SP_{Al}) and Iron (SP_{Fe})

In LSOM soils, added P accumulated mainly as SP_{Al}, was reported to be more available to plants than SP_{Fe}. The (control vs lime) and [(control + lime) vs LIOR] contrasts were not significant for $\Delta(SP_{Fe})$ (Table 10). The $r(SP_{Al})_F$ accounted for 79-86% of added P in LSOM soils, and for 35-58% in the HSOM soil (Fig. 6). The (control + lime) vs LIOR contrast was significant for $\Delta(SP_{Al})$ (Table 10). The LIOR decreased the proportions of SP_{Al} (Fig. 6). The $\Delta(SP_{Al})$ increased with added P ($P< 0.01$) (Fig. 6 and

Table 10) and was largest in LSOM soils. The $r(SP_{Fe})_F$ proportions were <11% for LSOM soils and 19-30% for the HSOM soil (Fig. 8).

Organic Phosphorus $(P_{org})_F$

The organic P turnover in soils is mediated by microbial activity and C dynamics (Huffman et al., 1996). In the highest P treatment, $\Delta(P_{org})$ increased with increasing organic C from 135-180 mg L^{-1} in control or limed LSOM soils, to 290-375 mg L^{-1} in LIOR-treated LSOM soils, 450-650 mg L^{-1} in control or limed HSOM, and 860 mg L^{-1} in LIOR-treated HSOM soils (Fig 8). The higher the OM content, the larger was $\Delta(P_{org})$. The (control vs lime) and [(control + lime) vs LIOR] contrasts were significant (Table 10). The higher the added P, the larger was the difference between treatments for $r(P_{org})_F$ (Fig. 9). For the highest P treatment in LSOM soils, $r(P_{org})_F$ was 1.8-2.4% for control, 1.7-4.6% for lime, and 3.8-5.0% for LIOR. In the HSOM soil, $r(P_{org})_F$ was 6.0% for control, 8.6% for lime, and 11.5% for LIOR (Table 11). Thus, some added P was converted into organic P in presence of lime or LIOR.

Desorbed Phosphorus Pool (DP)

The $\Delta(DP)$ increased abruptly with added P (Fig. 9). For the 5 P, 10 P, and 15 P treatments, respectively, $r(DP)_F$ increased from 3.2-7.8% to 10.1-13.2% and 17.6-22.6% in LSOM soils, and from 8.6-13.4% to 25.5-33.2% and 52.3-71.7% in the HSOM soil, respectively (Fig. 9). Comparatively, anion exchange resin P fractions ranged between 6% in high-P fixing soils and 74% in soils of low P sorption capacity across slightly to highly weathered soils of continental USA and Puerto Rico. The highest P treatment was the only one producing significant differences among control, lime, and LIOR treatments (Fig. 9). The $r(DP)_F$ averaged 21.5% in LSOM and 69.6% in HSOM soils. In the latter case, $r(DP)_F$ was 52.3% for the lime and 71.7% for the LIOR treatments. The organic ligands in LIOR presumably produced P desorption. The lime may increase or decrease P solubility depending on formation of new highly active polymeric hydroxy-Al, precipitation as insoluble Ca phosphates, or stimulation of microbial activities (Haynes 1982). In our case, microbial P immobilization rather than P sorption apparently decreased $\Delta(DP)$ by increasing $\Delta(P_{org})$ (Fig. 8).

Sorbed Phosphorus Pools

For the highest P treatment, $r(SP_{inorg})_F$ was found to be 78.9-88.4% for LSOM soils and 51.2-61.1% for the HSOM soil (Table 11). The $r(AP_{rev})_F$ averaged 16.5% in LSOM soils, and 36% in HSOM soil (Table 10). The $r(SP_{sr})_F$ accounted for 13.5% of the total sorbed P_i ($r(SP_{inorg})_F$) in the HSOM soil receiving LIOR compared to 61-79% for LSOM soils (Table 11).

The Phosphorus Partitioning in SOM Soil Groups

The flowchart in Fig. 10 illustrates the P pools for the highest P treatment in LSOM and HSOM soils receiving LIOR, as SOM was the single most determinant factor in reducing P retention in these soils. The $r(LBP)_F$ increased nearly 7 times from 5.0% in LSOM control soils to 34.0% in the HSOM soil receiving LIOR (Fig. 10), as total SOM increased about 3.3 times from 71 to 232 g kg^{-1}. The $r(DP)_F$ increased 3.3 times from 21.5% in LSOM control soils to 71.7 % in the HSOM soil receiving LIOR (Fig. 10). Conversely, $r(SP_{sr})_F$ decreased from 70.6% in LSOM control soils to 13.5% in the HSOM soil receiving LIOR. Added P was retained mainly as SP_{Al}, but P sorption varied among soils. The $r(SP_{Al})_F$ was 78.6% in LSOM soils without LIOR and 35.0% for the HSOM soil receiving LIOR (Fig. 7). Therefore, combining inorganic P and LIOR could improve P fertilizer efficiency in these high-P fixing soils. Since the effect of organic residues on P binding and desorption must depend not only on P but also on soil type (Olsen and Barber 1986), field trials are needed to ascertain the right proportions of LIOR and MAP maintaining maximum LBP in the fertilizer band during the potato growing season.

Results show that native (SOM) or supplemented (LIOR) sources of organic matter alleviated P fixation in podzolic soils, and modified the partition of added phosphate fertilizer (MAP) in favor of less tightly bound P pools. The LIOR appeared inefficient when applied at a rate of 23 g L^{-1} of soil to increase SOM by 14.6 g L^{-1} in soils already containing 40 to 50 mg SOM kg^{-1}. A comparative soil containing 200 mg SOM kg^{-1} and receiving similar amounts of LIOR in the same soil volume reduced the P fixation by Al from 78.6 to 35.0%, and increased loosely bound P from 5.0 to 34.0%. Since most acid mineral soils contain less than 50 g OM kg^{-1}, LIOR may improve fertilizer P efficiency in the fertilizer band of acid soils, thus potentially reducing P application rates for the potato production. A more detailed study is required to select the optimum organo-

mineral combinations for improving P efficiency in the fertilizer band in relation with amorphous and organically bound Al and Fe in acid light-textured soils used for the potato production. The P partitioning flowchart based on P fractionation indicated the major role of OM in reducing the P binding energy in those highly P-fixing soils.

Table 12

Effect of soil type and dry swine manure (LIOR) addition on Al- and Feextracted-pyrophosphate for the highest P treatment

		Pyrophosphate extraction (mmol kg^{-1})			
		Al_{pyr} [y]	Fe_{pyr} [y]	$(Al + Fe)_{pyr}$	$\dfrac{(Al + Fe)_{pyr}}{(Al + Fe)_{ox}} \times 100$
Soil	Treatment		Means		
Morin	No LIOR	80.0	32.1	112.1	19.6
	LIOR	81.5	32.1	113.6	19.9
Bevin	No LIOR	98.5	41.4	139.9	21.3
	LIOR	106.7	43.2	149.9	20.7
Ivry	No LIOR	55.6	20.0	75.6	16.5
	LIOR	78.5	20.7	99.2	18.0
Ivry peaty phase	No LIOR	35.6	22.9	58.4	24.5
	LIOR	39.3	25.7	65.0	27.2
	Df [z]		F value		
Soil	3	195.59**	906.54**	285.16**	97.96**
LIOR	1	21.63**	17.31**	24.43**	7.52*
LIOR x Soil	3	6.14*	3.77 ns	5.08*	3.98*
Root of error mean of squares		3.907	0.64	4.21	0.73
Coefficient of variation		5.4%	2.2%	4.1%	3.5%
R-Square		0.99	0.99	0.99	0.98

[z]df: degree of freedom; ns, *, **: non significant and significant at the 0.05 and the 0.01 levels, respectively;
[y].

EXAMPLE IV

DETERMINATION OF THE EFFECT OF THE COMPOSITION OF A LIOR-BASED OMF ON CORN GROWTH

To determine the effect of different combinations of biosolids and mineral fertilizers on the growth of corn, bio-treated manure was mixed to mineral fertilizers rich in N (32-0-0), P (8-25-3) or K(6-0-30), in different proportions. The parameters of the experimentation are described in Table 13 and Table 14.

Table 13

2001 characteristics of corn cultivated soils, corn cultural parameters and sample dates

Sites	1	2	3
Town or landlord	St-Madeleine	St-François	Montmagny
Producer	Francis Dion	Clément Lamonde	J. Yves Gosselin
Cultivar	Pioneer 38J54	Dkc27-11	Semico h12093
UTM	2800	2250	2300
Sowing date	May 3	May 18	May 19
Harvesting date	October 4	October 18	October 18
Plantlet sample date	June 15	June 27	June 27
Foliar sample date	August 2	August 8	
Soil sample date	May 25	October 18	October 18
Soil series	Richelieu	St-Épiphane	Kamouraska
Texture	Loam	Sandy loam	Slimy clay
Clay (%)	21.4	9.7	46.6
MO (%)	2.8	3.2	9.2
pH (0.01 M CaCl$_2$)	5.94	5.53	6.04
P$_{M-III}$(mg/kg)	95.8	48.4	89.9
Al$_{M-III}$(mg/kg)	790.8	426.2	1012.2
100(P/Al)$_{M-III}$(%)	12.1	11.4	8.9

Results

The effect of the different combinations of organo phosphatic fertilizer is shown in Table 14 and Fig. 11. Results show that 50 and 75% LIOR increases the yield of corn crops by 0.6 Tons/ha.

Table 14

Different combinaison effects of organo-mineral fertilizers with a LIOR basis on corn grain yield for three high phosphorus saturation sites (Montmagny, St-François, Ste-Madeleine)

Treatment	Yield	Grain Moisture	Density
kg P_2O_5/ha	Tons/ha	%	g/L
0 (Refefence)	8.213	34.58	776
20 (0% LIOR)	8.441	33.05	772
20 (25% LIOR)	8.594	34.22	776
20 (50% LIOR)	9.068	34.07	776
20 (75% LIOR)	9.065	34.33	776
20 (100% LIOR)	8.874	34.62	775
Error mean square	0.49	2.42	8.06
Variation factor (%)	5.47	7.08	1.04
			F Value
Site effect	141.90**	65.90*	16.14*
Treatment effect	4.21**	0.68ns	0.95ns
Block effect	4.86**	1.47ns	4.13*
Site*Block	8.52**	0.58ns	2.77*
Site*Treatment	1.03ns	0.95ns	0.86ns
Reference vs fertilized	10.85**	0.46ns	0.12ns
Linear effect (% LIOR)	6.54**	2.14ns	1.90ns
Quadratic effect (% LIOR)	3.55ns	0.27ns	2.64≈
Cubic effect (% LIOR)	0.95ns	0.38ns	0.02ns

EXAMPLE V

Determination on the effect of the composition of a DBM-based OMF on potatoe growth

To determine the effect of different combinations of biosolids and mineral fertilizers on the growth of potatoes, bio-treated manure was mixed to mineral fertilizers rich in N (32-0-0), P (8-25-3) or K(6-0-30), in different proportions. The parameters of the experimentation are described in Table 15.

Table 15

2001 characteristics of potatoes cultivated soils, cultural parameters and sample dates

Parameters	Sainte-Croix	Saint-Ubalde
Cultivar	GoldRush	GoldRush
Sowing date	May 21	May 17
Foliar sample date	July 13	July 18
Ridging date	July 6	July 11
Soil sample date	May 21	August 14
Harvesting date	September 18	September 8
Soil series	Tilly	Bevin
Texture	Slimy loam	Sandy loam
Clay (%)	21.0	6.5
MO (%)	3.9	4.9
pH (0.01 M CaCl$_2$)	5.10	5.35
Buffer pH	6.12	6.28
P_{M-III} (mg/kg)	21.6	79.2
Al_{M-III} (mg/kg)	1441.0	1760.5
$100(P/Al)_{M-III}$ (%)	1.5	4.5

Result

The effect of the different combinations of organo phosphatic fertilizer is shown in Table 16 and Fig. 12 and 13. Results show that 50% LIOR increases the yield of potatoes by approximately 6 tons/ha at both 75 and 150 kg P$_2$O$_5$/ha, which represents a 15% increase of the potatoes productivity.

Table 16

Dose and different combination of organo-mineral fertilizers with a LIOR basis effect on yield and specific weights of potatoes cultivation (cultivar Gold Rush, St-Ubalde)

Treatment		Yield	Specific weight	Tuber category > 57mm
kg P_2O_5/ha		Tons/ha		%
0 (Refefence)		35.079	1.075	70.0
75 (0% LIOR)		37.864	1.073	75.2
75 (25% LIOR)		39.505	1.072	76.4
75 (50% LIOR)		43.735	1.070	76.4
75 (75% LIOR)		40.625	1.070	78.7
75 (100% LIOR)		46.255	1.070	75.3
150 (0% LIOR)		40.433	1.074	76.0
150 (25% LIOR)		45.033	1.072	75.9
150 (50% LIOR)		46.420	1.073	78.0
150 (75% LIOR)		44.128	1.072	75.8
150 (100% LIOR)		48.728	1.069	75.3
Error mean square		4.40	0.004	4.06
Variation factor (%)		10.34	0.37	5.36
			F Value	
Treatment effect		3.56**	0.92ns	1.16ns
Block effect		5.88**	2.47≈	1.51ns
Reference	vs	12.62**	3.59≈	8.72**
fertilized				
Dose effect		7.93**	0.27ns	0.02ns
LInear dose		17.75**	2.77≈	7.76**
Quadratic dose		0.56ns	3.08≈	5.23**
Linear effect (% LIOR)		7.31**	3.16≈	0.02ns
Quadratic effect (% LIOR)		1.42ns	0.00ns	1.21ns
Cubic effect (% LIOR)		1.56ns	0.26ns	0.29ns

EXAMPLE VI

Determination of the effect of the composition of a LIOR-based OMF on soy growth

To determine the effect of different combinations of biosolids and mineral fertilizers on the growth of soy, bio-treated manure was mixed to mineral fertilizers rich in N (32-0-0), P (8-25-3) or K(6-0-30), in different proportions. The parameters of the experimentation are described in Table 17 and Table 18.

Table 17

2001 characteristics of soy cultivated soils, cultural parameters and sample dates

Parameters	Yvon Dion	Bernard Fontaine
Locality	St-Damase	St-Barnabé-Sud
Cultivar	Grand-Prix	Prograin Ohgata
UTM	2700	2625
Soil sample date	September 7	September 7
Harvesting date	September 7	September 7
Soil series	Ste-Rosalie	St-Hyacinthe
Texture	Sandy-argillaceous loam	Slimy loam
Clay (%)	20.9	17.8
MO (%)	2.1	2.1
pH (0.01 M CaCl$_2$)	5.58	5.62
buffer pH	7.04	7.06
P$_{M-III}$ (mg/kg)	157.2	106.1
Al$_{M-III}$ (mg/kg)	564.1	645.0
100(P/Al)$_{M-III}$ (%)	27.9	16.5

Table 18

2001 soy fertilization treatment at St-Damase and St-Barnabé South

Treatment	P Form	P Dose	MAP proportion	LIOR proportion	N	K$_2$O
		kg P$_2$O$_5$/ha		%	kg N/ha	kg K$_2$O/ha
A	(reference)	0	0	0	20	30
B	MAP	20	100	0	20	30
C	MAP+LIOR	20	75	25	20	30
D	MAP+LIOR	20	50	50	20	30
E	MAP+LIOR	20	25	75	20	30
F	LIORP	20	0	100	20	30

Results

The effect of the different combinations of organo phosphatic fertilizer is shown in Table 19 and Fig. 14. Results show that 75% LIOR increases the yield of soy by approximately 0.6 tons/ha, which represents a 21% increase of the soy productivity.

Table 19

Different combination of organo-mineral fertilizers with a LIOR basis effects on soy cultivation for the 2 sites (St-Barnabé South and St-Damase).

Treatment	Yield	Density
kg P$_2$O$_5$/ha	Tons/ha	g/L
0 (Refefence)	2.61	1.360
20 (0% LIOR)	2.85	1.381
20 (25% LIOR)	2.81	1.359
20 (50% LIOR)	3.47	1.364
20 (75% LIOR)	3.44	1.372
20 (100% LIOR)	3.18	1.365
Error mean square	0.49	0.02
Variation factor (%)	15.94	1.60
F Value		
Site effect	10.13**	0.07ns
Treatment effect	4.24**	1.16ns
Block effect	3.81**	0.23ns
Site*block	0.03ns	0.89ns
Site*treatment	0.67ns	0.14ns
Reference vs fertilized	7.63**	0.93ns
Linear effect (% LIOR)	3.61ten	0.67ns
Quadratic effect (% LIOR)	4.61*	1.38ns
Cubic effect (% LIOR)	3.92*	2.79tn

Experiments we conducted experiments since 1999 to determine the efficiency of a different OMF formula on the growth of different crops. OMF comprising 10 to 90% of LIOR were tested. The mineral fertilizer MAP (11-48-0) and bi-ammonium phosphate (18-46-0) were used to complete the OMF. Results demonstrated that an OMF comprising 80% (w/w) of BDM and 20% (w/w) of a mineral fertilizer is preferable.

The 80% BDM - 20% mineral fertilizer formula was tested on crops for a summer season. The 20% portion of mineral fertilizer comprised an equal amount of DAP and MAP. For example, 1 kg of BDM was supplemented with 145 grams of DAP and 145

gram of MAP. A commonly used binding agent have been added to the previous mixture (5% w/w of Min-U-Gel® 200 from Floridin, Quincy, FL, USA) to strengthen the bounds between every component of the pellet. Water have been added to the mixture to facilitate the homogenization, after which the mixture was extruded and dried for 24 hours.

Indeed, it respects the general rules regarding the chemical fertilizer relating to the nutrient composition since it comprise more than 24% of the combined macronutrient nitrogen (N), phosphorus (P_2O_5) and potassium (K). Moreover, the OMF of the present invention can provide an important amount of oligoelements required for the growth of crops (Cu, Zn, Boron, Molybdene, Manganese) since it is manure-based. The presence of binding agents, such as humic acids and fulvic acids, in the BDM represent a advantage of the OMF since they reversebly bind phosphorus, therefore facilitating its absorbtion by plant roots. The binding of phosphorus to humic or fulvic acids prevent its binding to iron or aluminum oxides, a process commonly observed in different types of soil.

While the invention has been described in connection with specific embodiments thereof, it will be understood that it is capable of further modifications and this application is intended to cover any variations, uses, or adaptations of the invention and including such departures from the present disclosure as come within known or customary practice within the art to which the invention pertains and as may be applied to the essential features hereinabove set forth, and as follows in the scope of the appended claims.